WHAT IS CLAIMED IS:

1. A compound having the formula (I):

$$\begin{array}{c|c}
Y_1^1 & X & Y_4^4 \\
Y_2^2 & Z & X_3^4 & R^1 \\
Z & X_3^4 & R^2 & R^2
\end{array}$$

$$\begin{array}{c|c}
R^4 & Q & L & R^3
\end{array}$$

4 wherein

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5 X is a member selected from the group consisting of a bond, -C(O)-,

6 $-C(R^5)(R^6)$ -, $-C(R^5)$ =, -S(O)-, $-S(O)_2$ - and -N=;

Z is a member selected from the group consisting of a bond, -N=, -O-, -S-,

 $-N(R^{17})$ - and $-C(R^{7})$ =, with the proviso that X and Z are not both a bond;

L is a member selected from the group consisting of a bond, C(O)-(C1-

C₈)alkylene, (C₁-C₈)alkylene and (C₂-C₈)heteroalkylene;

Q is a member selected from the group consisting of a bond, (C₁-

 $12 \qquad C_8) alkylene, (C_2-C_8) heteroalkylene, -C(O)-, -OC(O)-, -N(R^8)C(O)-, -CH_2CO-, -CH_2SO-ROC(O)-, -CH_2CO-, -CH_2CO-,$

and -CH2SO2-;

optionally L and Q can be linked together to form a 5- or 6-membered

heterocyclic group having from 1 to 3 heteroatoms;

 R^1 and R^2 are members independently selected from the group consisting of H, (C_1-C_8) alkyl, (C_2-C_8) heteroalkyl, aryl and heteroaryl, or optionally are combined to

18 form a 3 to 8-membered ring having from 0 to 2 heteroatoms as ring vertices;

optionally R² and L can be linked together to form a 5- or 6-membered

20 heterocyclic group having from 1 to 4 heteroatoms;

R3 is a member selected from the group consisting of hydroxy, (C1-

22 C₈)alkoxy, amino, (C₁-C₈)alkylamino, di(C₁-C₈)alkylamino, (C₂-C₈)heteroalkyl, (C₃-

Co)heterocyclyl, (C1-C8)acylamino, amidino, guanidino, ureido, cyano, heteroaryl.

24 -CONR⁹R¹⁰ and -CO₂R¹¹:

R4 is a member selected from the group consisting of (C1-C20)alkyl, (C2-

26 C₂₀)heteroalkyl, heteroaryl, aryl, heteroaryl(C₁-C₆)alkyl, heteroaryl(C₂-C₆)heteroalkyl,

27 arvl(C1-C6)alkvl and arvl(C2-C6)heteroalkvl;

R⁵ and R⁶ are each members independently selected from the group

29	consisting of H, (C ₁ -C ₈)alkyl, (C ₂ -C ₈)heteroalkyl, heteroaryl and aryl, or optionally R ⁵
30	and R ⁶ are combined to form a 3- to 7-membered ring;
31	R ⁷ and R ⁸ are each members independently selected from the group
32	consisting of H, (C1-C8)alkyl, (C2-C8)heteroalkyl, heteroaryl and aryl,
33	each R ⁹ , R ¹⁰ and R ¹¹ is independently selected from the group consisting
34	of H, (C_1-C_8) alkyl, (C_2-C_8) heteroalkyl, heteroaryl, aryl, heteroaryl (C_1-C_6) alkyl,
35	$heteroaryl(C_2\text{-}C_8)heteroalkyl,\ aryl(C_1\text{-}C_8)alkyl\ and\ aryl(C_2\text{-}C_8)heteroalkyl;$
36	Y ¹ and Y ² are each members independently selected from the group
37	consisting of $-C(R^{12})=$, $-N=$, $-O-$, $-S-$ and $-N(R^{13})-$;
38	Y ³ is a member selected from the group consisting of N and C wherein the
39	carbon atom shares a double bond with either Z or Y4; and
40	Y^4 is a member selected from the group consisting of -N(R ¹⁴)-, -C(R ¹⁴)=,
41	-N= and -N(R^{14})-C(R^{15})(R^{16})-, wherein
42	each R ¹² is a member independently selected from the group consisting of
43	$H,halogen,hydroxy,amino,alkylamino,dialkylamino,(C_1-C_8)alkyl,(C_2-C_8)heteroalkyl,$
44	heteroaryl and aryl, or optionally when Y^1 and Y^2 are both $-C(R^{12})=$ the two R^{12} groups
45	can be combined to form a substituted or unsubstituted 5- to 6-membered cycloalkyl,
46	heterocycloalkyl, aryl or heteroaryl ring; or optionally when Y^1 is $-C(R^{12})$ = and X is $-$
47	$C(R^5)$ = or $-C(R^5)(R^6)$ -, R^{12} and R^5 can be combined to form a substituted or unsubstituted
48	5- to 6-membered cycloalkyl, heterocycloalkyl, aryl or heteroaryl ring;
49	R ¹³ is a member selected from the group consisting of H, (C ₁ -C ₈)alkyl,
50	$(C_2\text{-}C_8) \\ \text{heteroaryl}, \text{ aryl, heteroaryl} (C_1\text{-}C_6) \\ \text{alkyl, heteroaryl} (C_2\text{-}C_8) \\ \text{heteroalkyl, heteroaryl} \\ \text{aryl, heteroaryl} (C_1\text{-}C_6) \\ \text{alkyl, heteroaryl} (C_2\text{-}C_8) \\ \text{heteroaryl} ($
51	$aryl(C_1-C_8)alkyl$ and $aryl(C_2-C_8)heteroalkyl$;
52	R ¹⁴ is a member selected from the group consisting of (C ₁ -C ₈)alkyl, (C ₂ -
53	$C_8) heteroalkyl, aryl(C_1 - C_8) alkyl, aryl(C_2 - C_8) heteroalkyl, heteroaryl(C_1 - C_8) alkyl, aryl(C_2 - C_8) heteroalkyl, heteroaryl(C_1 - C_8) alkyl, heteroalkyl, heteroalkyl,$
54	heteroaryl(C2-C8)heteroalkyl, heteroaryl and aryl;
55	$R^{15} \ \text{and} \ R^{16}$ are each members independently selected from the group
56	consisting of H, (C1-C8)alkyl and (C2-C8)heteroalkyl; and
57	R ¹⁷ is a member selected from the group consisting of H, (C ₁ -C ₈)alkyl,
58	$(C_2\text{-}C_8) \\ \text{heteroaryl}, \\ \text{aryl}, \\ \text{heteroaryl}(C_1\text{-}C_6) \\ \text{alkyl}, \\ \text{heteroaryl}(C_2\text{-}C_8) \\ \text{heteroalkyl}, \\ \text{heteroaryl}(C_2\text{-}C_8) \\ \text{heteroalkyl}, \\ \text{heteroaryl}(C_2\text{-}C_8) \\ \text{heteroaryl}(C_2\text{-}$
59	$\operatorname{aryl}(C_1-C_8)$ alkyl and $\operatorname{aryl}(C_2-C_8)$ heteroalkyl, or optionally when Y^2 is $-C(R^{12})=$ or $-$
60	$N(R^{13})$ -, R^{17} can be combined with R^{12} or R^{13} to form a substituted or unsubstituted 5- to
61	6-membered cycloalkyl, heterocycloalkyl, aryl or heteroaryl ring;
62	with the proviso that when the Y ³ -containing ring system is a

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- 63 quinazolinone or quinolinone ring system, and R⁴-Q- is substituted or unsubstituted (C₅-
- 64 C₁₅)alkyl, then R³-L- is other than substituted or unsubstituted (C₂-C₈)alkylene or a
- 65 substituted or unsubstituted (C2-C8)heteroalkylene attached to -NR'R", wherein R' and
- 66 R" are independently selected from the group consisting of hydrogen and (C1-C8)alkyl, or
- 67 optionally are combined with the nitrogen atom to which each is attached to form a 5-, 6-
- 68 or 7-membered ring.
 - A compound of Claim 1, wherein Y⁴ is -N(R¹⁴)- wherein R¹⁴ is
 selected from the group consisting of aryl and heteroaryl.
 - A compound of Claim 1, wherein X is -C(O)-
 - A compound of Claim 1, wherein Z is -N=.
 - 5. A compound of Claim 1, wherein Y^1 and Y^2 are each $-C(R^{12})=$
 - wherein the two R¹² groups are combined to form a fused 6-membered aryl or heteroaryl
 ring.
 - 6. A compound of Claim 1, wherein X is -C(O)-; Z is -N=; Y^3 is C; and Y^1 and Y^2 are each $-C(R^{12})=$.
 - 7. A compound of Claim 6, wherein the two R¹² groups are combined to
 form a fused 6-membered substituted or unsubstituted aryl or heteroaryl ring.
 - 8. A compound of Claim 6, wherein Y⁴ is -N(R¹⁴)-.
 - A compound of Claim 6, wherein Y⁴ is -C(R¹⁴)=.
 - 1 10. A compound of Claim 7, wherein Y⁴ is -N(R¹⁴)-.
 - A compound of Claim 7, wherein Y⁴ is -C(R¹⁴)=.
 - 1 12. A compound of Claim 1, wherein L is (C₁-C₈)alkylene.
 - 1 13. A compound of Claim 1, wherein Q is -C(O)-.
 - 14. A compound of Claim 1, wherein R⁴ is selected from the group
 consisting of (C₂-C₁₅)alkyl, substituted or unsubstituted phenyl and biphenyl.

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 Y^1 and Y^2 are each $-C(R^{12})=$.

15. A compound of Claim 1, wherein R³ is selected from the group 1 consisting of (C1-C8)alkoxy, (C1-C8)alkylamino, di(C1-C8)alkylamino, (C2-2 C8) heteroalkyl, (C3-C9) heterocyclyl, (C1-C8) acylamino, cyano, heteroaryl, -CONR 9R10 3 and -CO2R11. 4 16. A compound of Claim 1, wherein R¹ and R² are independently selected 1 from the group consisting of H and (C1-C4)alkyl. 2 17. A compound of Claim 1, wherein Y³ is C and the carbon atom shares a 1 2 double bond with Z. 18. A compound of Claim 1, wherein X is $-C(R^5)(R^6)$ -; Y^4 is $-N(R^{14})$ -. 1 wherein R¹⁴ is substituted or unsubstituted arvl or heteroaryl; Y³ is C; Z is -N=; and Y¹ 2 and Y2 are each -C(R12)=. 3 19. A compound of Claim 18, wherein X is -CH₂- and the R¹² groups are 1 combined to form a substituted or unsubstituted aryl or heteroaryl ring. 2 20. A compound of Claim 1, wherein X is $-C(R^5)=$; Y^4 is $-C(R^{14})=$. 1 wherein R¹⁴ is substituted or unsubstituted aryl or heteroaryl; Y³ is C; Z is -N=; and Y¹ 2 and Y2 are each -C(R12)=. 3 21. A compound of Claim 20, wherein R1 is H. 1 22. A compound of Claim 1, wherein X is a bond; Y4 is -N(R14)-, wherein 1 R¹⁴ is substituted or unsubstituted aryl or heteroaryl; Y³ is C; Z is -N=; and Y¹ and Y² are 2 each -C(R12)=. 3 23. A compound of Claim 22, wherein the R12 groups are combined to 1 2 form a substituted or unsubstituted aryl or heteroaryl ring. 24. A compound of Claim 22, wherein R1 is H. 1

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wherein R¹⁴ is substituted or unsubstituted arvl or heteroarvl; Y³ is C; Z is -C(R⁷)=; and

25. A compound of Claim 1, wherein X is -C(R⁵)=; Y⁴ is -C(R¹⁴)=,

26. A compound of Claim 25, wherein R⁵ and R¹² are combined to form a 1 5- or 6-membered substituted or unsubstituted aryl or heteroaryl ring. 2 27. A compound of Claim 25, wherein R1 is H. 1 28. A compound of Claim 1, wherein X is a bond: Z is -N= or $-N(R^{17})$ -: 1 Y⁴ is -C(R¹⁴)=, wherein R¹⁴ is substituted or unsubstituted aryl or heteroaryl; Y¹ is 2 selected from the group consisting of -O-, -S- and $-N(R^{13})$ -; and Y^2 is $-C(R^{12})$ =. 3 29. A compound of Claim 28, wherein Y1 is -O- and Z is -N=. 1 30. A compound of Claim 28, wherein Y1 is -S- and Z is -N=. 1 31. A compound of Claim 28, wherein Y^1 is $-N(R^{13})$ - and Z is -N=. 1 32. A compound of Claim 1, wherein X is -SO₂-; Y⁴ is -N(R¹⁴)=, wherein R^{14} is substituted or unsubstituted arvl or heteroaryl; Y^3 is C; Z is -N= or $-C(R^7)=$; and Y^1 2 and Y^2 are each $-C(R^{12})=$. 3 33. A compound of Claim 32, wherein R1 is H. 1 34. A compound of Claim 1, wherein X is a bond; Z is -O-, -S- or 1 $-N(R^{17})$ -; Y^1 is -N= or $-N(R^{13})$ -; Y^2 is $-C(R^{12})$ =; and Y^4 is $-C(R^{14})$ = wherein R^{14} is 2 substituted or unsubstituted aryl or heteroaryl. 3 35. A compound of Claim 34, wherein Y1 is -N= and Z is -O-. 1 36. A compound of Claim 34, wherein Y^1 is -N= and Z is -S-. 1 37. A compound of Claim 34, wherein Z is -N(R¹⁷)-. 1 38. A compound of Claim 34, wherein R1 is H. 1 39. A compound of Claim 1, wherein X is a bond; Y1 is -N(R13)- or =N-; 1 Y^2 is $-C(R^{12})=: Y^3$ is $C: Y^4$ is $-C(R^{14})=$ wherein R^{14} is substituted or unsubstituted aryl or 2 heteroaryl; and Z is $-N(R^{17})$ - or =N-, with the proviso that Y^1 and Z are not both =N-. 3 40. A compound of Claim 1, wherein X is a bond; Y1 and Y2 are each 1 independently -C(R¹²)=: Y³ is C: Y⁴ is -C(R¹⁴)= wherein R¹⁴ is substituted or 2

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- unsubstituted aryl or heteroaryl; and Z is $-N(R^{17})$ -, O or S. 3 41. A compound of Claim 40, wherein the two R¹² groups are combined to 1 form a fused 5- or 6-membered substituted or unsubstituted aryl or heteroaryl ring. 2 42. A compound of Claim 1, wherein X is -C(O)-; Y¹ is -N(R¹³)-; Y² is 1 -N=: Y³ is C; Y⁴ is -N(R¹⁴)- wherein R¹⁴ is substituted or unsubstituted aryl or heteroaryl; 2 3 and Z is a bond. 43. A compound of Claim 42, wherein R1 is H. 1 44. A compound of Claim 1, wherein X is-C(O)-; Z is -N(R¹⁷)- wherein 1 R¹⁷ is substituted or unsubstituted aryl or heteroaryl; Y¹ and Y² are each independently 2 $-C(R^{12})=: Y^3 \text{ is C: and } Y^4 \text{ is -N}=.$ 3 45. A compound of Claim 44, wherein R1 is H. 1 46. A compound of Claim 1, wherein X and Z are -N=, Y1 and Y2 are each 1 independently $-C(R^{12})=: Y^3$ is C: and Y^4 is $-C(R^{14})=$ wherein R^{14} is a substituted or 2 unsubstituted arvl or heteroarvl group. 3 47. A compound of Claim 46, wherein R1 is H. 1 48. A compound of Claim 1, wherein X is -C(O)-; Y⁴ is 1 -N(R¹⁴)-C(R⁵)(R⁶)-; wherein R¹⁴ is substituted or unsubstituted aryl or heteroaryl; Y¹ and 2 Y^2 are each independently $-C(R^{12})=$; Y^3 is C; and Z is -N=. 3 49. A compound of Claim 48. wherein R1 is H. 1 50. A compound of Claim 1, wherein the Y³-containing ring system is 1
 - pyridine, pyrazine and benzodiazepine.

selected from the group consisting of quinoline, quinazoline, naphthalene, quinolinone,

quinazolinone, triazolinone, pyrimidin-4-one, benzimidazole, thiazole, imidazole,

51. A compound of Claim 1, having the formula (III):

$$(R_a)_n$$
 $X N$ R^1 R^4 Q N L R^2

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A4 is C or N: 5

6 X is -CO-, -CH2- or a bond;

R1 and R2 are each members independently selected from the group consisting of 7 H and (C1-C4)alkyl; 8

> R¹⁴ is a substituted or unsubstituted member selected from the group consisting of phenyl, pyridyl, thiazolyl, thienyl and pyrimidinyl;

O is -CO-;

L is (C1-C8)alkylene;

the subscript n is an integer of from 0 to 4; and

each Ra is independently selected from the group consisting of halogen, -OR',

-OC(O)R', -NR'R", -SR', -R', -CN, -NO2, -CO2R', -CONR'R", -C(O)R',

-OC(O)NR'R", -NR"C(O)R', -NR"C(O)2R', ,-NR'-C(O)NR"R"',

-NH-C(NH2)=NH, -NR'C(NH2)=NH, -NH-C(NH2)=NR', -S(O)R', -

S(O)₂R', -S(O)₂NR'R", -N₃, -CH(Ph)₂, perfluoro(C₁-C₄)alkoxy, and

perfluoro(C1-C4)alkyl, wherein R', R" and R" are each independently selected from the group consisting of H, (C1-C8)alkyl, (C2-C8)heteroalkyl,

unsubstituted aryl, unsubstituted heteroaryl, (unsubstituted aryl)-(C1-

C₄)alkyl, and (unsubstituted aryl)oxy-(C₁-C₄)alkyl.

52. A compound of Claim 51, wherein X is -C(O)-.

53. A compound of Claim 51, wherein X is -CH2-.

54. A compound of Claim 51, wherein X is a bond.

55. A compound of Claim 51, wherein R⁴ is substituted or unsubstituted benzyl, wherein said substituents are selected from the group consisting of halogen, 2

halo(C1-C4)alkyl, halo(C1-C4)alkoxy, cyano, nitro, and phenyl.

1	56. A compound of Claim 51, wherein R ¹⁴ is selected from the group
2	consisting of substituted phenyl, substituted pyridyl, substituted thiazolyl and substituted
3	thienyl, wherein the substituents are selected from the group consisting of cyano, halogen,
4	(C1-C8)alkoxy, (C1-C8)alkyl, (C2-C8)heteroalkyl, CONH2, methylenedioxy and
5	ethylenedioxy.
1	57. A compound of Claim 51, wherein R ¹⁴ is substituted phenyl, wherein
2	the substituents are selected from the group consisting of cyano, halogen, (C1-C8)alkoxy,
3	$(C_1\hbox{-} C_8)$ alkyl, $(C_2\hbox{-} C_8)$ heteroalkyl, CONH2, methylenedioxy and ethylenedioxy.
1	58. A compound of Claim 51, wherein R ⁴ is substituted or unsubstituted
2	benzyl, wherein said substituents are selected from the group consisting of halogen,
3	halo(C1-C4)alkyl, halo(C1-C4)alkoxy, cyano, nitro and phenyl, and R14 is substituted
4	phenyl, wherein the substituents are selected from the group consisting of cyano, halogen,
5	(C1-C8)alkoxy, (C1-C8)alkyl, (C2-C8)heteroalkyl, CONH2, methylenedioxy and
6	ethylenedioxy.
1	59. A compound of Claim 51, wherein R1 is selected from the group
2	consisting of methyl, ethyl and propyl, and R ² is hydrogen.
1	60. A compound of Claim 51, wherein \mathbb{R}^1 and \mathbb{R}^2 are each methyl.
1	61. A compound of Claim 51, wherein R3 is selected from the group
2	consisting of (C1-C8)alkoxy, amino, (C1-C8)alkylamino, di(C1-C8)alkylamino, (C2-
3	C ₈)heteroalkyl, (C ₃ -C ₉)heterocyclyl and heteroaryl.
1	62. A compound of Claim 51, wherein R ³ is selected from the group
2	consisting of substituted or unsubstituted pyridyl and substituted or unsubstituted
3	imidazolyl.
1	63. A compound of Claim 51, wherein L is (C_1-C_4) alkylene.
1	64. A compound of Claim 51, wherein X is -CO-; R1 and R2 are each
2	independently selected from the group consisting of H, methyl and ethyl; R14 is phenyl;
3	L is methylene, ethylene or propylene, R ³ is selected from the group consisting of
4	substituted or unsubstituted pyridyl and substituted or unsubstituted imidazolyl; R4 is
5	substituted or unsubstituted benzyl, wherein said substituents are selected from the group

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- 6 consisting of halogen, halo(C1-C4)alkyl, halo(C1-C4)alkoxy, cyano, nitro, and phenyl; and
- 7 each R_a is selected from the group consisting of halogen, -OR', -OC(O)R', -NR'R", -SR',
- 8 -R', -CN, -NO₂, -CO₂R', -CONR'R", -C(O)R', -NR"C(O)R', -NR'-C(O)NR"R"',
- 9 perfluoro(C1-C4)alkoxy, and perfluoro(C1-C4)alkyl, wherein R', R" and R" are each
- 10 independently selected from the group consisting of H, (C1-C8)alkyl, (C2-C8)heteroalkyl,
- 11 unsubstituted aryl, unsubstituted heteroaryl, (unsubstituted aryl)-(C1-C4)alkyl, and
- 12 (unsubstituted aryl)oxy-(C₁-C₄)alkyl.
 - 65. A compound of Claim 51, wherein said compound is selected from the group consisting of:

66. A pharmaceutical composition comprising a pharmaceutically acceptable carrier or excipient and a compound having the formula (I):

5 wherein

X is a member selected from the group consisting of a bond, -C(O)-, -C(R⁵)(R⁶)-, -C(R⁵)=, -S(O)-, -S(O)₂- and -N=;

8	Z is a member selected from the group consisting of a bond, $-N=$, $-O-$, $-S-$,
9	$-N(R^{17})$ - and $-C(R^7)$ =, with the proviso that X and Z are not both a bond;
10	L is a member selected from the group consisting of a bond, C(O)-(C1-
11	C ₈)alkylene, (C ₁ -C ₈)alkylene and (C ₂ -C ₈)heteroalkylene;
12	Q is a member selected from the group consisting of a bond, (C1-
13	$C_8) alkylene, (C_2\text{-}C_8) heteroalkylene, -C(O), -OC(O), -N(R^8)C(O), -CH_2CO, -CH_2SO$
14	and -CH ₂ SO ₂ -;
15	optionally L and Q can be linked together to form a 5- or 6-membered
16	heterocyclic group having from 1 to 3 heteroatoms;
17	R ¹ and R ² are members independently selected from the group consisting
18	of H, (C1-C8)alkyl, (C2-C8)heteroalkyl, aryl and heteroaryl, or optionally are combined to
19	form a 3 to 8-membered ring having from 0 to 2 heteroatoms as ring vertices;
20	optionally R ² and L can be linked together to form a 5- or 6-membered
21	heterocyclic group having from 1 to 4 heteroatoms;
22	R ³ is a member selected from the group consisting of hydroxy, (C ₁ -
23	$C_8) alkoxy, amino, (C_1-C_8) alkylamino, di (C_1-C_8) alkylamino, (C_2-C_8) heteroalkyl, (C_3-C_8) alkylamino, di (C_1-C_8) alkylamino, di (C_1$
24	C ₉)heterocyclyl, (C ₁ -C ₈)acylamino, amidino, guanidino, ureido, cyano, heteroaryl,
25	-CONR 9 R 10 and -CO $_{2}$ R 11 ;
26	R ⁴ is a member selected from the group consisting of (C ₁ -C ₂₀)alkyl, (C ₂ -
27	$C_{20}) heteroalkyl,\ heteroaryl,\ aryl,\ heteroaryl(C_1-C_6) alkyl,\ heteroaryl(C_2-C_6) heteroalkyl,$
28	$aryl(C_1-C_6)alkyl$ and $aryl(C_2-C_6)heteroalkyl$;
29	R ⁵ and R ⁶ are each members independently selected from the group
30	consisting of H, (C_1-C_8) alkyl, (C_2-C_8) heteroalkyl, heteroaryl and aryl, or optionally R^5
31	and R ⁶ are combined to form a 3- to 7-membered ring;
32	R ⁷ and R ⁸ are each members independently selected from the group
33	consisting of H, (C1-C8)alkyl, (C2-C8)heteroalkyl, heteroaryl and aryl,
34	each R^9,R^{10} and R^{11} is independently selected from the group consisting
35	of H, (C_1-C_8) alkyl, (C_2-C_8) heteroalkyl, heteroaryl, aryl, heteroaryl (C_1-C_6) alkyl,
36	$heteroaryl (C_2-C_8) heteroalkyl, aryl (C_1-C_8) alkyl \ and \ aryl (C_2-C_8) heteroalkyl;$
37	Y1 and Y2 are each members independently selected from the group
38	consisting of $-C(R^{12})=$, $-N=$, $-O-$, $-S-$ and $-N(R^{13})-$;
39	Y3 is a member selected from the group consisting of N and C wherein the
40	carbon atom shares a double bond with either Z or Y4; and
41	Y^4 is a member selected from the group consisting of -N(R^{14})-, -C(R^{14})=,

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-N= and -N(R14)-C(R15)(R16)-, wherein
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                       each R12 is a member independently selected from the group consisting of
43
       H. halogen, hydroxy, amino, alkylamino, dialkylamino, (C1-C8)alkyl, (C2-C8)heteroalkyl,
44
       heteroaryl and aryl, or optionally when Y1 and Y2 are both -C(R12)= the two R12 groups
45
       can be combined to form a substituted or unsubstituted 5- to 6-membered cycloalkyl,
46
       heterocycloalkyl, aryl or heteroaryl ring; or optionally when Y1 is -C(R12)= and X is -
47
       C(R^5)= or -C(R^5)(R^6)-, R^{12} and R^5 can be combined to form a substituted or unsubstituted
48
       5- to 6-membered cycloalkyl, heterocycloalkyl, aryl or heteroaryl ring;
49
                        R13 is a member selected from the group consisting of H, (C1-C8)alkyl,
50
       (C2-C8)heteroalkyl, heteroaryl, aryl, heteroaryl(C1-C6)alkyl, heteroaryl(C2-C8)heteroalkyl,
51
       aryl(C1-C8)alkyl and aryl(C2-C8)heteroalkyl;
52
                        R<sup>14</sup> is a member selected from the group consisting of (C<sub>1</sub>-C<sub>8</sub>)alkyl, (C<sub>2</sub>-
53
       C_8)heteroalkyl, aryl(C_1-C_8)alkyl, aryl(C_2-C_8)heteroalkyl, heteroaryl(C_1-C_8)alkyl,
54
       heteroaryl(C2-C8)heteroalkyl, heteroaryl and aryl;
55
                        R<sup>15</sup> and R<sup>16</sup> are each members independently selected from the group
56
       consisting of H, (C1-C8)alkyl and (C2-C8)heteroalkyl; and
57
                        R<sup>17</sup> is a member selected from the group consisting of H, (C<sub>1</sub>-C<sub>8</sub>)alkyl,
58
       (C2-C8)heteroalkyl, heteroaryl, aryl, heteroaryl(C1-C6)alkyl, heteroaryl(C2-C8)heteroalkyl,
59
       aryl(C<sub>1</sub>-C<sub>8</sub>)alkyl and aryl(C<sub>2</sub>-C<sub>8</sub>)heteroalkyl, or optionally when Y<sup>2</sup> is -C(R<sup>12</sup>)= or -
60
       N(R<sup>13</sup>)-, R<sup>17</sup> can be combined with R<sup>12</sup> or R<sup>13</sup> to form a substituted or unsubstituted 5- to
61
       6-membered cycloalkyl, heterocycloalkyl, aryl or heteroaryl ring;
62
                        with the proviso that when the Y<sup>3</sup>-containing ring system is a
63
       quinazolinone or quinolinone ring system, and R4-Q- is substituted or unsubstituted (C5-
64
       C<sub>15</sub>)alkyl, then R<sup>3</sup>-L- is other than substituted or unsubstituted (C<sub>2</sub>-C<sub>8</sub>)alkylene or a
65
       substituted or unsubstituted (C2-C8)heteroalkylene attached to -NR'R", wherein R' and
66
       R" are independently selected from the group consisting of hydrogen and (C1-C8)alkyl, or
67
       optionally are combined with the nitrogen atom to which each is attached to form a 5-, 6-
68
69
       or 7-membered ring.
                        67. A composition of Claim 66, wherein Y<sup>4</sup> is -N(R<sup>14</sup>)- wherein R<sup>14</sup> is
  1
        selected from the group consisting of aryl and heteroaryl.
  2
  1
                        68. A composition of Claim 66, wherein X is -C(O)-.
```

69. A composition of Claim 66, wherein Z is -N=.

1	70. A composition of Claim 66, wherein Y^1 and Y^2 are each $-C(R^{12})=$		
2	wherein the two R ¹² groups are combined to form a fused 6-membered aryl or heteroary		
3	ring.		
l	71. A composition of Claim 66, wherein X is -C(O)-; Z is -N=; Y³ is C;		
2	and Y1 and Y2 are each -C(R12)= wherein the two R12 groups are combined to form a		
3	fused 6-membered substituted or unsubstituted aryl or heteroaryl ring.		
1	72. A composition of Claim 66, wherein L is $(C_1\text{-}C_8)$ alkylene.		
1	73. A composition of Claim 66, wherein Q is -C(O)		
1	74. A composition of Claim 66, wherein R ⁴ is selected from the group		
2	consisting of $(C_5\text{-}C_{15})$ alkyl, substituted or unsubstituted phenyl and biphenyl.		
1	75. A composition of Claim 66, wherein R ³ is selected from the group		
2	consisting of (C1-C8)alkoxy, (C1-C8)alkylamino, di(C1-C8)alkylamino, (C2-		
3	C ₈)heteroalkyl, (C ₃ -C ₉)heterocyclyl, (C ₁ -C ₈)acylamino, cyano, heteroaryl, -CONR ⁹ R ¹⁰		
4	and $-CO_2R^{11}$.		
1	76. A composition of Claim 66, wherein \mathbb{R}^1 and \mathbb{R}^2 are independently		
2	selected from the group consisting of H and (C ₁ -C ₄)alkyl.		
1	77. A composition of Claim 66, wherein Y ³ is C and the carbon atom		
2	shares a double bond with Z.		
1	78. A composition of Claim 66, wherein the Y³-containing ring system is		
2	selected from the group consisting of quinoline, quinazoline, naphthalene, quinolinone,		
3	quinazolinone, triazolinone, pyrimidin-4-one, benzimidazole, thiazole, imidazole,		
4	pyridine, pyrazine and benzodiazepine.		
1	79. A composition of Claim 66, wherein the compound has the formula		
2	(III):		
	(R ₂) - R ¹⁴		

4	Ш
5	wherein
6	A^4 is C or N;
7	X is -CO-, -CH ₂ - or a bond;
8	R1 and R2 are each members independently selected from the group
9	consisting of H and (C ₁ -C ₄)alkyl;
.0	R ¹⁴ is a substituted or unsubstituted member selected from the group
1	consisting of phenyl, pyridyl, thiazolyl, thienyl and pyrimidinyl;
2	Q is -CO-;
3	L is (C ₁ -C ₈)alkylene;
4	the subscript n is an integer of from 0 to 4; and
5	each Ra is independently selected from the group consisting of halogen, -
16	OR', -OC(O)R', -NR'R", -SR', -R', -CN, -NO ₂ , -CO ₂ R', -CONR'R", -C(O)R',
17	$-OC(O)NR'R", -NR"C(O)R', -NR"C(O)_2R', ,-NR'-C(O)NR"R"", -NH-C(NH_2) = NH, -NR^2 - C(O)NR^2 - NH - C(NH_2) = NH - C(NH_2) + $
18	$NR'C(NH_2) = NH, -NH-C(NH_2) = NR', -S(O)R', -S(O)_2R', -S(O)_2NR'R'', -N_3, -CH(Ph)_2,$
19	perfluoro(C_1 - C_4)alkoxy, and perfluoro(C_1 - C_4)alkyl, wherein R', R" and R" are each
20	independently selected from the group consisting of, (C_1-C_8) alkyl, (C_2-C_8) heteroalkyl,
21	unsubstituted aryl, unsubstituted heteroaryl, (unsubstituted aryl)-(C_1 - C_4)alkyl, and
22	(unsubstituted aryl)oxy-(C ₁ -C ₄)alkyl.
1	80. A composition in accordance with Claim 79, wherein X is $-C(O)$
1	81. A composition in accordance with Claim 79, wherein X is $-CH_2$
1	82. A composition in accordance with Claim 79, wherein X is a bond.
1	83. A composition in accordance with Claim 79, wherein R4 is substituted
2	or unsubstituted benzyl, wherein said substituents are selected from the group consisting
3	of halogen, halo (C_1-C_4) alkyl, halo (C_1-C_4) alkoxy, cyano, nitro, and phenyl.
1	84. A composition in accordance with Claim 79, wherein R ¹⁴ is selected
2	from the group consisting of substituted phenyl, substituted pyridyl, substituted thiazolyl
3	and substituted thienyl, wherein the substituents are selected from the group consisting of
4	cvano, halogen, (C ₁ -C ₈)alkoxy, (C ₁ -C ₈)alkyl, (C ₂ -C ₈)heteroalkyl, CONH ₂ ,

methylenedioxy and ethylenedioxy.

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aryl)oxy-(C1-C4)alkyl.

1	85. A composition in accordance with Claim 79, wherein \mathbb{R}^1 is selected
2	from the group consisting of methyl, ethyl and propyl, and R ² is.
1	86. A composition in accordance with Claim 79, wherein \mathbb{R}^1 and \mathbb{R}^2 are
2	each methyl.
1	87. A composition in accordance with Claim 79, wherein R ³ is selected
2	from the group consisting of substituted or unsubstituted pyridyl and substituted or
3	unsubstituted imidazolyl.
1	88. A composition in accordance with Claim 79, wherein L is (C1-
2	C ₄)alkylene.
1	89. A composition in accordance with Claim 79, wherein X is -CO-; R ¹
2	and R ² are each independently selected from the group consisting of, methyl and ethyl;
3	R^{14} is selected from the group consisting of substituted or unsubstituted phenyl; L is
4	methylene, ethylene or propylene, R ³ is selected from the group consisting of substituted
5	or unsubstituted pyridyl and substituted or unsubstituted imidazolyl; R4 is substituted or
6	unsubstituted benzyl, wherein said substituents are selected from the group consisting of
7	halogen, halo(C_1 - C_4)alkyl, halo(C_1 - C_4)alkoxy, cyano, nitro, and phenyl; and each R_a is
8	selected from the group consisting of halogen, -OR', -OC(O)R', -NR'R", -SR', -R', -CN,
9	$-NO_2, -CO_2R', -CONR'R'', -C(O)R', -NR''C(O)R', -NR''-C(O)NR''R''', perfluoro(C_1-CO)R''R''', -C(O)R''R''', -C(O)R'', -NR''C(O)R'', -NR''-C(O)R'', -NR''', -C(O)R'', -C(O)R''$
10	C ₄)alkoxy, and perfluoro(C ₁ -C ₄)alkyl, wherein R', R" and R" are each independently
11	selected from the group consisting of, (C1-C8)alkyl, (C2-C8)heteroalkyl, unsubstituted

90. The composition of Claim 79, wherein said compound is:

aryl, unsubstituted heteroaryl, (unsubstituted aryl)- (C_1-C_4) alkyl, and (unsubstituted

1 2 3

91. A method of treating an inflammatory or immune condition or disease in a subject, said method comprising administering to a subject in need of such treatment a therapeutically effective amount of a compound having the formula (I):

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16

wherein

X is a member selected from the group consisting of a bond, -C(O)-,

 $-C(R^5)(R^6)$ -, $-C(R^5)$ =, -S(O)-, $-S(O)_2$ - and -N=; 8

Z is a member selected from the group consisting of a bond, -N=, -O-, -S-,

 $-N(R^{17})$ - and $-C(R^7)$ =, with the proviso that X and Z are not both a bond;

L is a member selected from the group consisting of a bond, C(O)-(C1-

C₈)alkylene, (C₁-C₈)alkylene and (C₂-C₈)heteroalkylene; 12

O is a member selected from the group consisting of a bond, (C1-

 C_8)alkylene, (C_2-C_8) heteroalkylene, -C(O)-, -OC(O)-, $-N(R^8)C(O)$ -, $-CH_2CO$ -, $-CH_2SO$ -14

and -CH2SO2-:

optionally L and Q can be linked together to form a 5- or 6-membered

heterocyclic group having from 1 to 3 heteroatoms; 17

18	R ¹ and R ² are members independently selected from the group consisting
19	of H, (C1-C8)alkyl, (C2-C8)heteroalkyl, aryl and heteroaryl, or optionally are combined to
20	form a 3 to 8-membered ring having from 0 to 2 heteroatoms as ring vertices;
21	optionally R ² and L can be linked together to form a 5- or 6-membered
22	heterocyclic group having from 1 to 4 heteroatoms;
23	R ³ is a member selected from the group consisting of hydroxy, (C ₁ -
24	C_8)alkoxy, amino, (C_1-C_8) alkylamino, di (C_1-C_8) alkylamino, (C_2-C_8) heteroalkyl, (C_3-C_8) alkylamino, di (C_1-C_8) alkylamino, di $($
25	C ₉)heterocyclyl, (C ₁ -C ₈)acylamino, amidino, guanidino, ureido, cyano, heteroaryl,
26	-CONR 9 R 10 and -CO ₂ R 11 ;
27	R4 is a member selected from the group consisting of (C1-C20)alkyl, (C2-
28	$C_{20}) heteroalkyl,\ heteroaryl,\ aryl,\ heteroaryl(C_1-C_6) alkyl,\ heteroaryl(C_2-C_6) heteroalkyl,$
29	aryl(C1-C6)alkyl and aryl(C2-C6)heteroalkyl;
30	R ⁵ and R ⁶ are each members independently selected from the group
31	consisting of H, (C_1-C_8) alkyl, (C_2-C_8) heteroalkyl, heteroaryl and aryl, or optionally \mathbb{R}^5
32	and R ⁶ are combined to form a 3- to 7-membered ring;
33	R ⁷ and R ⁸ are each members independently selected from the group
34	consisting of H, (C1-C8)alkyl, (C2-C8)heteroalkyl, heteroaryl and aryl,
35	each R ⁹ , R ¹⁰ and R ¹¹ is independently selected from the group consisting
36	of H, (C_1-C_8) alkyl, (C_2-C_8) heteroalkyl, heteroaryl, aryl, heteroaryl (C_1-C_6) alkyl,
37	$heteroaryl(C_2\text{-}C_8) heteroalkyl, aryl(C_1\text{-}C_8) alkyl and aryl(C_2\text{-}C_8) heteroalkyl;$
38	Y1 and Y2 are each members independently selected from the group
39	consisting of $-C(R^{12})=$, $-N=$, $-O-$, $-S-$ and $-N(R^{13})-$;
40	Y³ is a member selected from the group consisting of N and C wherein the
41	carbon atom shares a double bond with either Z or Y4; and
42	Y^4 is a member selected from the group consisting of $-N(R^{14})$ -, $-C(R^{14})$ =,
43	$-N=$ and $-N(R^{14})-C(R^{15})(R^{16})-$, wherein
44	each R ¹² is a member independently selected from the group consisting of
45	H, halogen, hydroxy, amino, alkylamino, dialkylamino, (C1-C8)alkyl, (C2-C8)heteroalkyl,
46	heteroaryl and aryl, or optionally when Y^1 and Y^2 are both $-C(R^{12})$ = the two R^{12} groups
47	can be combined to form a substituted or unsubstituted 5- to 6-membered cycloalkyl,
48	heterocycloalkyl, aryl or heteroaryl ring; or optionally when Y^1 is $-C(R^{12})$ = and X is $-$
49	$C(R^5)$ = or $-C(R^5)(R^6)$ -, R^{12} and R^5 can be combined to form a substituted or unsubstituted
50	5- to 6-membered cycloalkyl, heterocycloalkyl, aryl or heteroaryl ring;
51	R ¹³ is a member selected from the group consisting of H, (C ₁ -C ₈)alkyl,

7

 $(C_2\text{-}C_8) \\ \text{heteroaryl}, \text{ aryl, heteroaryl} \\ (C_1\text{-}C_6) \\ \text{alkyl, heteroaryl} \\ (C_2\text{-}C_8) \\ \text{heteroalkyl, heteroaryl} \\ (C_2\text{-}C_8) \\ \text{heteroaryl} \\ (C_2\text{-}C_8) \\ (C_2\text{-}C$ 52 aryl(C1-C8)alkyl and aryl(C2-C8)heteroalkyl; 53 R¹⁴ is a member selected from the group consisting of (C₁-C₈)alkyl, (C₂-54 C_8)heteroalkyl, aryl(C_1 - C_8)alkyl, aryl(C_2 - C_8)heteroalkyl, heteroaryl(C_1 - C_8)alkyl, 55 heteroaryl(C2-C8)heteroalkyl, heteroaryl and aryl; 56 R¹⁵ and R¹⁶ are each members independently selected from the group 57 consisting of H, (C1-C8)alkyl and (C2-C8)heteroalkyl; and 58 R¹⁷ is a member selected from the group consisting of H, (C₁-C₈)alkyl, 59 (C_2-C_8) heteroalkyl, heteroaryl, aryl, heteroaryl (C_1-C_6) alkyl, heteroaryl (C_2-C_8) heteroalkyl, 60 $arvl(C_1-C_8)$ alkyl and $arvl(C_2-C_8)$ heteroalkyl, or optionally when Y^2 is $-C(R^{12})$ = or -61 $N(R^{13})$ -, R^{17} can be combined with R^{12} or R^{13} to form a substituted or unsubstituted 5- to 62 6-membered cycloalkyl, heterocycloalkyl, aryl or heteroaryl ring; 63 with the proviso that when the Y3-containing ring system is a 64 quinazolinone or quinolinone ring system, and R4-Q- is substituted or unsubstituted (C5-65 C15)alkyl, then R3-L- is other than substituted or unsubstituted (C2-C8)alkylene or a 66 substituted or unsubstituted (C2-C8)heteroalkylene attached to -NR'R", wherein R' and 67 R" are independently selected from the group consisting of hydrogen and (C1-C8)alkyl, or 68 optionally are combined with the nitrogen atom to which each is attached to form a 5-, 6-69 or 7-membered ring. 70 92. The method of Claim 91, wherein said compound is administered 1 2 orally, parenterally or topically. 93. The method of Claim 91, wherein said compound modulates CXCR3. 1 94. The method of Claim 91, wherein said compound is a CXCR3 1 2 antagonist. 95. The method of Claim 91, wherein said inflammatory or immune 1 condition or disease is selected from the group consisting of neurodegenerative diseases, 2 multiple sclerosis, systemic lupus erythematosus, rheumatoid arthritis, atherosclerosis, 3 encephalitis, meningitis, hepatitis, nephritis, sepsis, sarcoidosis, psoriasis, eczema, 4 uticaria, type I diabetes, asthma, conjunctivitis, otitis, allergic rhinitis, chronic obstructive 5

pulmonary disease, sinusitis, dermatitis, inflammatory bowel disease, ulcerative colitis,

2.

8 organ transplant conditions and skin transplant conditions.

96. The method of Claim 91, wherein said compound is administered in combination with a second therapeutic agent, wherein said second therapeutic agent is useful for treating or preventing neurodegenerative diseases, multiple sclerosis, systemic lupus erythematosus, rheumatoid arthritis, atherosclerosis, encephalitis, meningitis, hepatitis, nephritis, sepsis, sarcoidosis, psoriasis, eczema, uticaria, type I diabetes, asthma, conjunctivitis, otitis, allergic rhinitis, chronic obstructive pulmonary disease, sinusitis, dermatitis, inflammatory bowel disease, ulcerative colitis, Crohn's disease, Behcet's syndrome, gout, cancer, viral infections, bacterial infections, organ transplant conditions or skin transplant conditions.

97. A method of treating a CXCR3-mediated condition or disease in a subject, said method comprising administering to a subject in need of such treatment a therapeutically effective amount of a compound having the formula (I):

wherein

X is a member selected from the group consisting of a bond, -C(O)-, -C(R 5)(R 6)-, -C(R 5)=, -S(O)-, -S(O)₂- and -N=;

 $Z \ is \ a \ member \ selected \ from \ the \ group \ consisting \ of \ a \ bond, -N=, -O-, -S-, -N(R^{17})- \ and -C(R^7)=, \ with \ the \ proviso \ that \ X \ and \ Z \ are \ not \ both \ a \ bond;$

 $L \ is \ a \ member \ selected \ from \ the \ group \ consisting \ of \ a \ bond, \ C(O)-(C_1-C_8) alkylene, \ (C_1-C_8) alkylene \ and \ (C_2-C_8) heteroalkylene;$

Q is a member selected from the group consisting of a bond, (C_1-C_8) alkylene, (C_2-C_8) heteroalkylene, -C(O), -OC(O), $-N(R^8)C(O)$, $-CH_2CO$, $-CH_2SO$ and $-CH_2SO_2$;

optionally L and Q can be linked together to form a 5- or 6-membered heterocyclic group having from 1 to 3 heteroatoms;

R¹ and R² are members independently selected from the group consisting

19	of H, (C_1-C_8) alkyl, (C_2-C_8) heteroalkyl, aryl and heteroaryl, or optionally are combined to
20	form a 3 to 8-membered ring having from 0 to 2 heteroatoms as ring vertices;
21	optionally R ² and L can be linked together to form a 5- or 6-membered
22	heterocyclic group having from 1 to 4 heteroatoms;
23	R ³ is a member selected from the group consisting of hydroxy, (C ₁ -
24	C ₈)alkoxy, amino, (C ₁ -C ₈)alkylamino, di(C ₁ -C ₈)alkylamino, (C ₂ -C ₈)heteroalkyl, (C ₃ -
25	C ₉)heterocyclyl, (C ₁ -C ₈)acylamino, amidino, guanidino, ureido, cyano, heteroaryl,
26	-CONR ⁹ R ¹⁰ and -CO ₂ R ¹¹ ;
27	R ⁴ is a member selected from the group consisting of (C ₁ -C ₂₀)alkyl, (C ₂ -
28	$C_{20}) heteroalkyl,\ heteroaryl,\ aryl,\ heteroaryl(C_1-C_6) alkyl,\ heteroaryl(C_2-C_6) heteroalkyl,$
29	$aryl(C_1-C_6)alkyl$ and $aryl(C_2-C_6)heteroalkyl$;
30	R ⁵ and R ⁶ are each members independently selected from the group
31	consisting of H, (C1-C8)alkyl, (C2-C8)heteroalkyl, heteroaryl and aryl, or optionally R5
32	and R ⁶ are combined to form a 3- to 7-membered ring;
33	R ⁷ and R ⁸ are each members independently selected from the group
34	consisting of H, (C1-C8)alkyl, (C2-C8)heteroalkyl, heteroaryl and aryl,
35	each R ⁹ , R ¹⁰ and R ¹¹ is independently selected from the group consisting
36	of H, (C1-C8)alkyl, (C2-C8)heteroalkyl, heteroaryl, aryl, heteroaryl(C1-C6)alkyl,
37	heteroaryl(C2-C8)heteroalkyl, aryl(C1-C8)alkyl and aryl(C2-C8)heteroalkyl;
38	Y ¹ and Y ² are each members independently selected from the group
39	consisting of $-C(R^{12})=$, $-N=$, $-O-$, $-S-$ and $-N(R^{13})-$;
40	Y3 is a member selected from the group consisting of N and C wherein the
41	carbon atom shares a double bond with either Z or Y4; and
42	Y^4 is a member selected from the group consisting of -N(R ¹⁴)-, -C(R ¹⁴)=,
43	$-N=$ and $-N(R^{14})-C(R^{15})(R^{16})-$, wherein
44	each R ¹² is a member independently selected from the group consisting of
45	H, halogen, hydroxy, amino, alkylamino, dialkylamino, (C1-C8)alkyl, (C2-C8)heteroalkyl,

-N= and -N(R¹⁴)-C(R¹⁵)(R¹⁶)-, wherein each R¹² is a member independently selected from the group consisting of H, halogen, hydroxy, amino, alkylamino, dialkylamino, (C₁-C₈)alkyl, (C₂-C₈)heteroalkyl, heteroaryl and aryl, or optionally when Y¹ and Y² are both -C(R¹²)= the two R¹² groups can be combined to form a substituted or unsubstituted 5- to 6-membered cycloalkyl, heterocycloalkyl, aryl or heteroaryl ring; or optionally when Y¹ is -C(R¹²)= and X is -C(R⁵)= or -C(R⁵)(R⁶)-, R¹² and R⁵ can be combined to form a substituted or unsubstituted 5- to 6-membered cycloalkyl, heterocycloalkyl, aryl or heteroaryl ring;

R¹³ is a member selected from the group consisting of H, (C₁-C₈)alkyl,

 $(C_2\text{-}C_8) \\ \text{heteroalkyl, heteroaryl, aryl, heteroaryl} \\ (C_1\text{-}C_6) \\ \text{alkyl, heteroaryl} \\ (C_2\text{-}C_8) \\ \text{heteroalkyl, heteroaryl} \\ (C_2\text{-}C_8) \\ \text{heteroalkyl, heteroaryl} \\ \text{heteroaryl} \\$

C₈)alkylene.

53	aryl(C1-C8)alkyl and aryl(C2-C8)heteroalkyl;
54	R ¹⁴ is a member selected from the group consisting of (C ₁ -C ₈)alkyl, (C ₂ -
55	$C_8) heteroalkyl, aryl(C_1-C_8) alkyl, aryl(C_2-C_8) heteroalkyl, heteroaryl(C_1-C_8) alkyl, heteroa$
56	heteroaryl(C2-C8)heteroalkyl, heteroaryl and aryl;
57	R ¹⁵ and R ¹⁶ are each members independently selected from the group
58	consisting of H, (C1-C8)alkyl and (C2-C8)heteroalkyl; and
59	R ¹⁷ is a member selected from the group consisting of H, (C ₁ -C ₈)alkyl,
60	$(C_2-C_8) \\ heteroaryl, \ aryl, \ heteroaryl \\ (C_1-C_6) \\ alkyl, \ heteroaryl \\ (C_2-C_8) \\ heteroalkyl, \ heteroaryl \\ (C_3-C_8) \\ heteroalkyl, \ heteroaryl \\ heteroaryl $
61	$aryl(C_1-C_8)alkyl$ and $aryl(C_2-C_8)heteroalkyl$, or optionally when Y^2 is $-C(R^{12})=$ or $-$
62	N(R ¹³)-, R ¹⁷ can be combined with R ¹² or R ¹³ to form a substituted or unsubstituted 5- to
63	6-membered cycloalkyl, heterocycloalkyl, aryl or heteroaryl ring;
64	with the proviso that when the Y3-containing ring system is a
65	quinazolinone or quinolinone ring system, and R4-Q- is substituted or unsubstituted (C5-
66	C ₁₅)alkyl, then R ³ -L- is other than substituted or unsubstituted (C ₂ -C ₈)alkylene or a
67	substituted or unsubstituted (C2-C8)heteroalkylene attached to -NR'R", wherein R' and
68	R" are independently selected from the group consisting of hydrogen and (C1-C8)alkyl, or
69	optionally are combined with the nitrogen atom to which each is attached to form a 5-, 6-
70	or 7-membered ring.
	98. A method in accordance with Claim 97, wherein Y^4 is $-N(R^{14})$ -
1	wherein R ¹⁴ is selected from the group consisting of aryl and heteroaryl.
2	wherein R** is selected from the group consisting of aryl and neteroaryl.
1	99. A method in accordance with Claim 97, wherein X is -C(O)
1	100. A method in accordance with Claim 97, wherein Z is $-N=$.
1	101. A method in accordance with Claim 97, wherein Y ¹ and Y ² are
2	each -C(R ¹²)=, wherein the two R ¹² groups are combined to form a fused 6-membered
3	aryl or heteroaryl ring.
1	102. A method in accordance with Claim 97, wherein X is -C(O)-; Z is
2	$-N=$; Y^3 is C; and Y^1 and Y^2 are each $-C(R^{12})=$ wherein the two R^{12} groups are combined
3	to form a fused 6-membered substituted or unsubstituted aryl or heteroaryl ring.
1	103. A method in accordance with Claim 97, wherein L is (C ₁ -

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1	104.	A method in accordance with Claim 97, wherein Q is $-C(O)$
1	105.	A method in accordance with Claim 97, wherein R4 is selected
2	from the group consis	sting of (C ₅ -C ₁₅)alkyl, substituted or unsubstituted phenyl and
3	biphenyl.	
	106	A method in accordance with Claim 97, wherein R ³ is selected
1	106.	A method in accordance with Claim y , wherein X is selected sting of (C_1-C_8) alkya, (C_1-C_8) alkylamino, di (C_1-C_8) alkylamino,
2		
3		C ₃ -C ₉)heterocyclyl, (C ₁ -C ₈)acylamino, cyano, heteroaryl,
4	-CONR ⁹ R ¹⁰ and -CC	$\rho_2 R^{++}$.
1	107.	A method in accordance with Claim 97, wherein R ¹ and R ² are
2	independently selecte	ed from the group consisting of H and (C ₁ -C ₄)alkyl.
1	108.	A method in accordance with Claim 97, wherein Y ³ is C and the
2	carbon atom shares a	double bond with Z.
1	109.	A method in accordance with Claim 97, wherein the Y3-containing
2	ring system is selecte	ed from the group consisting of quinoline, quinazoline, naphthalene,
3		linone, triazolinone, pyrimidin-4-one, benzimidazole, thiazole,
4	-	pyrazine and benzodiazepine.
	-	
1	110.	A method in accordance with Claim 97, wherein said compound
2	has the formula (III)	:
		$(R_a)_n \stackrel{\wedge}{\underset{\wedge}{\sqcup}} A^4 \stackrel{\wedge}{\underset{\wedge}{\vee}} R^{14}$
3		R ⁴ -Q ^{-N} -L-R ³
4		Ш
5	wherein	
6	A4 is C or N;	
7	X is -CO-, -	CH ₂ - or a bond;
8	R1 and R2 are	e each members independently selected from the group consisting of
0	TI on	1 (C, -C,)alkyl:

phenyl, pyridyl, thiazolyl, thienyl and pyrimidinyl;

R¹⁴ is a substituted or unsubstituted member selected from the group consisting of

1

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12
             O is -CO-;
             L is (C<sub>1</sub>-C<sub>8</sub>)alkylene;
13
             the subscript n is an integer of from 0 to 4; and
14
             each Ra is independently selected from the group consisting of halogen, -OR',
15
                     -OC(O)R', -NR'R", -SR', -R', -CN, -NO2, -CO2R', -CONR'R", -C(O)R',
16
                     -OC(O)NR'R", -NR"C(O)R', -NR"C(O)2R', ,-NR'-C(O)NR"R"",
17
                     -NH-C(NH<sub>2</sub>)=NH, -NR'C(NH<sub>2</sub>)=NH, -NH-C(NH<sub>2</sub>)=NR', -S(O)R', -
18
                     S(O)_2R', -S(O)_2NR'R'', -N_3, -CH(Ph)_2, perfluoro(C_1-C_4)alkoxy, and
19
                     perfluoro(C1-C4)alkyl, wherein R', R" and R" are each independently
20
                     selected from the group consisting of H, (C1-C8)alkyl, (C2-C8)heteroalkyl,
21
                      unsubstituted aryl, unsubstituted heteroaryl, (unsubstituted aryl)-(C1-
22
                      C<sub>4</sub>)alkyl, and (unsubstituted aryl)oxy-(C<sub>1</sub>-C<sub>4</sub>)alkyl.
23
                             A method in accordance with Claim 110, wherein X is -C(O)-.
 1
                      111.
                             A method in accordance with Claim 110, wherein X is -CH2-.
                      112.
 1
                             A method in accordance with Claim 110, wherein X is a bond.
                      113.
                             A method in accordance with Claim 110, wherein R4 is substituted
 1
                      114.
      or unsubstituted benzyl, wherein said substituents are selected from the group consisting
 2
      of halogen, halo(C1-C4)alkyl, halo(C1-C4)alkoxy, cyano, nitro, and phenyl.
 3
                             A method in accordance with Claim 110, wherein R14 is selected
 1
      from the group consisting of substituted phenyl, substituted pyridyl, substituted thiazolyl
 2
      and substituted thienyl, wherein the substituents are selected from the group consisting of
 3
      cyano, halogen, (C1-C8)alkoxy, (C1-C8)alkyl, (C2-C8)heteroalkyl, CONH2,
 4
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A method in accordance with Claim 110, wherein R1 is selected 116. from the group consisting of methyl, ethyl and propyl, and R² is hydrogen. 2

methylenedioxy and ethylenedioxy.

- A method in accordance with Claim 110, wherein R1 and R2 are 1 117. each methyl. 2
- A method in accordance with Claim 110, wherein R3 is selected 1 from the group consisting of substituted or unsubstituted pyridyl and substituted or 2

2

- 3 unsubstituted imidazolyl.
- 1 119. A method in accordance with Claim 110, wherein L is (C₁-
- 2 C₄)alkylene.
- 1 120. A method in accordance with Claim 110, wherein X is -CO-; R¹
- 2 and R² are each independently selected from the group consisting of H, methyl and ethyl;
- $3 \quad R^{14}$ is selected from the group consisting of substituted or unsubstituted phenyl; Q is –
- 4 CO-; L is methylene, ethylene or propylene, R³ is selected from the group consisting of
- 5 substituted or unsubstituted pyridyl and substituted or unsubstituted imidazolyl; R4 is
- 6 substituted or unsubstituted benzyl, wherein said substituents are selected from the group
- 7 consisting of halogen, halo(C₁-C₄)alkyl, halo(C₁-C₄)alkoxy, cyano, nitro, and phenyl; and
- 8 each R_a is selected from the group consisting of halogen, -OR', -OC(O)R', -NR'R", -SR',
- 9 -R', -CN, -NO₂, -CO₂R', -CONR'R", -C(O)R', -NR"C(O)R', -NR'-C(O)NR"R",
- 10 perfluoro(C₁-C₄)alkoxy, and perfluoro(C₁-C₄)alkyl, wherein R', R" and R" are each
- 11 independently selected from the group consisting of H, (C₁-C₈)alkyl, (C₂-C₈)heteroalkyl,
- 12 unsubstituted aryl, unsubstituted heteroaryl, (unsubstituted aryl)-(C1-C4)alkyl, and
- 13 (unsubstituted aryl)oxy-(C₁-C₄)alkyl.
 - 121. The method of Claim 110, wherein said compound is selected from the group consisting of:

1	122. A method in accordance with Claim 97, wherein said CXCR3-
2	mediated condition is selected from the group consisting of neurodegenerative diseases,
3	multiple sclerosis, systemic lupus erythematosus, rheumatoid arthritis, atherosclerosis,
4	encephalitis, meningitis, hepatitis, nephritis, sepsis, sarcoidosis, psoriasis, eczema,
5	uticaria, type I diabetes, asthma, conjunctivitis, otitis, allergic rhinitis, chronic obstructive
6	pulmonary disease, sinusitis, dermatitis, inflammatory bowel disease, ulcerative colitis,
7	Crohn's disease, Behcet's syndrome, gout, cancer, viral infections, bacterial infections,
8	organ transplant conditions and skin transplant conditions.

- 1 123. The method of Claim 97, wherein said compound modulates 2 CXCR3.
 - 124. A method in accordance with Claim 110, wherein said compound is administered in combination with a second therapeutic agent, wherein said second therapeutic agent is useful for treating neurodegenerative diseases, multiple sclerosis, systemic lupus erythematosus, rheumatoid arthritis, atherosclerosis, encephalitis, meningitis, hepatitis, nephritis, sepsis, sarcoidosis, psoriasis, eczema, uticaria, type I diabetes, asthma, conjunctivitis, otitis, allergic rhinitis, chronic obstructive pulmonary disease, sinusitis, dermatitis, inflammatory bowel disease, ulcerative colitis, Crohn's disease, Behcet's syndrome, gout, cancer, viral infections, bacterial infections, organ transplant conditions or skin transplant conditions.
 - 125. A method in accordance with Claim 124, wherein said organ transplant condition is a bone marrow transplant condition or a solid organ transplant condition.
 - 126. A method in accordance with Claim 125, wherein said solid organ transplant condition is a kidney transplant condition, a liver transplant condition, a lung transplant condition, a heart transplant condition or a pancreas transplant condition.
 - 127. A method in accordance with Claim 97, wherein said CXCR3mediated condition is restenosis.
 - 128. A method in accordance with Claim 97, wherein said CXCR3mediated condition is selected from the group consisting of multiple sclerosis, rheumatoid

1

2

3	arthritis and organ transplant conditions.
1	129. A method in accordance with Claim 110, wherein said compound
2	is used in conjunction with another therapeutic agent selected from the group consisting
3	of Remicade®, Enbrel®, a COX-2 inhibitor, a glucocorticoid, an immunosuppressant,
4	methotrexate, predisolone, azathioprine, cyclophosphamide, tacrolimus, mycophenolate,
5	hydroxychloroquine, sulfasalazine, cyclosporine A, D-penicillamine, a gold compound,
6	an antilymphocyte or antithymocyte globulin, betaseron, avonex and copaxone.
1	130. A method in accordance with Claim 110, wherein said CXCR3-
2	mediated condition is an organ transplant condition and said compound is used alone or i
3	combination with a second therapeutic agent selected from the group consisting of
4	cyclosporine A, FK-506, rapamycin, mycophenolate, prednisolone, azathioprene,
5	cyclophosphamide and an antilymphocyte globulin.
1	131. A method in accordance with Claim 110, wherein said CXCR3-
2	mediated condition is rheumatoid arthritis and said compound is used alone or in
3	combination with a second therapeutic agent selected from the group consisting of
4	methotrexate, sulfasalazine, hydroxychloroquine, cyclosporine A, D-penicillamine,
5	Remicade®, Enbrel®, auranofin and aurothioglucose.
1	132. A method in accordance with Claim 110, wherein said CXCR3-
2	mediated condition is multiple sclerosis and said compound is used alone or in
3	combination with a second therapeutic agent selected from the group consisting of
4	betaseron, avonex, azathioprene, capoxone, prednisolone and cyclophosphamide.
1	133. The method of Claim 110, wherein said subject is a human.
1	134. A method for the modulation of CXCR3 function in a cell,

135. A method for the modulation of CXCR3 function, comprising contacting a CXCR3 protein with a compound of Claim 1.

comprising contacting said cell with a compound of Claim 1.